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<p>To aid in the development of boron-enhanced fluorinitramino explosives, the kinetics of individual reactions are measured over wide temperature ranges. It is found that the pre-exponentials of rate coefficients of BO reactions tend to be two orders of magnitude smaller than for the A1O reactions with the same oxidants. This may explain why no reaction could be observed between BO and CO<sub>2</sub>,  <math>k &lt; 1 \times 10^{-14} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}</math>.</p>				
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## PROGRESS

### Introduction

The goals of this program are:

- (i) to make accurate measurements, over wide temperature ranges, of rate coefficients for boron combustion in C/H/N/O/F environments, needed for models used in the development of new underwater explosives, and
- (ii) to use the measurements to obtain a further understanding, to allow predictions for additional reactions occurring with such explosives.

To this end measurements are made in unique high temperature reactors of the HTFFR (high-temperature fast-flow reactor) and HTP (high-temperature photochemistry) type. These are to be supplemented, where warranted, by semi-empirical and *ab initio* studies.

### Results

A series of measurements have now been made on BO reactions. The remarkable observation is that the pre-exponentials of the rate coefficients are much lower than for the corresponding AlO reactions. Witness the following two pairs, where the rate coefficients are expressed in  $\text{cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$  units:

- (1)  $\text{BO} + \text{O}_2 \rightarrow \text{BO}_2 + \text{O}$ ;  $k(300\text{-}1000 \text{ K}) = 7.9 \times 10^{-12} \exp(161 \text{ K/T})$  (Ref. 1)
- (2)  $\text{AlO} + \text{O}_2 \rightarrow \text{AlO}_2 + \text{O}$ ;  $k(1200\text{-}1690 \text{ K}) = 7.7 \times 10^{-10} \exp(-10008 \text{ K/T})$  (Ref. 2)
- (3)  $\text{BO} + \text{HCl} \rightarrow (\text{HBO} + \text{Cl})$  and/or  $(\text{OBCl} + \text{H})$ ; (Ref. 1)  
 $k(300\text{-}760 \text{ K}) = 6.3 \times 10^{-13} \exp(-1403 \text{ K/T})$
- (4)  $\text{AlO} + \text{HCl} \rightarrow (\text{HAlO} + \text{Cl})$  and/or  $(\text{OAlCl} + \text{H})$ ; (Ref. 3)  
 $k(440\text{-}1590 \text{ K}) = 5.6 \times 10^{-11} \exp(-139 \text{ K/T})$

The exothermic  $\text{BO} + \text{CO}_2 \rightarrow \text{BO}_2 + \text{CO}$  reaction (5) was found to be immeasurably slow,<sup>1</sup>  $k(300\text{-}1200 \text{ K}) < 3 \times 10^{-14}$ . The  $\text{AlO} + \text{CO}_2$  reaction, proceeded with  $k$  about  $4 \times 10^{-14}$  in this temperature range.<sup>4</sup> An *ab initio* study of reaction (5) showed evidence for a  $\text{BOCO}_2$  adduct, but no barrier.<sup>5</sup> Hence, a small pre-exponential should be the cause of the small  $k(5)$ . Current boron combustion models are very sensitive to this reaction.

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## PLANS

Further experimental studies planned include the reactions of BO with N<sub>2</sub>O, and HF, and of BF with BF<sub>3</sub>, O<sub>2</sub> and H<sub>2</sub>O.

The reason for the above-noted differences in the pre-exponential parts of the rate coefficients of the monoxides of the two group 13 elements B and Al is being considered. In previous work from this laboratory much smaller differences between BCl and AlCl rate coefficients were observed.<sup>6,7,8</sup> The BCl reactions with O<sub>2</sub> and CO<sub>2</sub> are actually somewhat faster than their AlCl counterparts. For N<sub>2</sub>O the reverse relation holds. Semi-empirical SECI theory was applied successfully to predict the relation between the exponential terms for a series of those reactions.<sup>9</sup> This approach might be extended to the monoxides. These combined considerations are significant for allowing predictions of  $k(T)$  values, in addition to those measured, with greater accuracy than appears in the current literature for such refractory species reactions.

## PARTICIPANTS AND CONTACTS

The principal participant on this project was J.-D.R. Rocha, an MS/Ph.D. student.

## PUBLICATIONS AND AWARDS

1. D.P. Belyung, G.T. Dalakos, Q. Zhang, J.D.-R. Rocha, and Fontijn A., "Wide Temperature Range Studies of BO and BO<sub>2</sub> Reactions", Abstract for The Fourth International Conference on Chemical Kinetics at NIST, July 1997. A full-length paper by the same title is in preparation.

There were no Awards this year.

## REFERENCES

1. D.P. Belyung, G.T. Dalakos, Q. Zhang, J.D.-R. Rocha, and A. Fontijn, "Wide Temperature Range Studies of BO and BO<sub>2</sub> Reactions", Abstract for The Fourth International Conference on Chemical Kinetics, NIST, Gaithersburg, MD, July 1997.
2. D.P. Belyung and A. Fontijn, "The AlO + O<sub>2</sub> Reaction System over a Wide Temperature Range", J. Phys. Chem. 99, 12225 (1995).
3. A.G. Slavejkov, C.T. Stanton, and A. Fontijn, "High-Temperature Fast-Flow Reactor Kinetics Studies of the Reactions of AlO with Cl<sub>2</sub> and HCl over Wide-Temperature Ranges", J. Phys. Chem. 94, 3347 (1990).
4. D.F. Rogowski, A.J. English, and A. Fontijn, "A High-Temperature Fast-Flow Reactor Kinetics Study of the Reaction AlO + CO<sub>2</sub> → AlO<sub>2</sub> + CO. Thermochemical Implications" J. Phys. Chem. 90, 1688 (1986).
5. P. Marshall, University of North Texas, Private Communication.

6. A.G. Slavejko, P.M. Futerko, and A. Fontijn, "High-Temperature Fast-Flow Reactor Kinetics Study of the Reaction Between BCl and CO<sub>2</sub> from 770 to 1830 K" *Twenty-Third Symposium (International) on Combustion*, (The Combustion Institute, Pittsburgh, 1990), p. 155
7. P.M. Futerko and A. Fontijn, "Experimental and Transition-State Theory Studies of the Gas-Phase Reactions of AlCl with N<sub>2</sub>O, CO<sub>2</sub>, and SO<sub>2</sub>", *J. Phys. Chem.* 97, 7222 (1993).
8. P.M. Futerko, A.G. Slavejko, and A. Fontijn, "Wide Temperature Range Kinetics of the Gas-Phase Reactions of BCl with SO<sub>2</sub>, N<sub>2</sub>O, O<sub>2</sub>, and CO<sub>2</sub>", *J. Phys. Chem.* 97, 11950 (1993).
9. A.S. Blue, D.P. Belyung, and A. Fontijn, "Activation Barriers for Series of Exothermic Homologous Reactions. V. Boron Group Diatomic Species Reactions", *J. Chem. Phys.* 107, 3791 (1997).